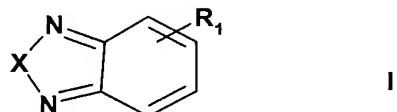


**Amendments to the Claims:**

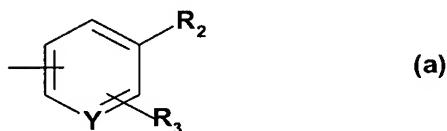
This listing of claims will replace all prior versions and listings of claims in the application:

**Listing of Claims:**

1. (Original) A compound of formula I



wherein X is O or S, R<sub>1</sub> is 5-(2-fluoro-ethylamino)-thiazol-2-yl, 5-(2-<sup>18</sup>F-ethylamino)-thiazol-2-yl or a group of formula (a)



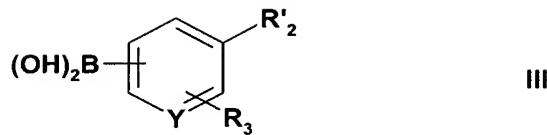
wherein Y is CH or N, R<sub>2</sub> is NHCH<sub>3</sub>, NH<sup>11</sup>CH<sub>3</sub>, N(CH<sub>3</sub>)<sup>11</sup>CH<sub>3</sub>, N(CH<sub>3</sub>)<sub>2</sub>, N(<sup>11</sup>CH<sub>3</sub>)<sub>2</sub>, NH(CH<sub>2</sub>)<sub>n</sub>F, NH(CH<sub>2</sub>)<sub>n</sub><sup>18</sup>F, N(CH<sub>3</sub>)-(CH<sub>2</sub>)<sub>n</sub>F, N(CH<sub>3</sub>)-(CH<sub>2</sub>)<sub>n</sub><sup>18</sup>F, O-(CH<sub>2</sub>)<sub>n</sub>F, O-(CH<sub>2</sub>)<sub>n</sub><sup>18</sup>F, CONH(CH<sub>2</sub>)<sub>n</sub>F or CONH(CH<sub>2</sub>)<sub>n</sub><sup>18</sup>F (n being in each case 2 to 4) and R<sub>3</sub> is hydroxy, (C1-4)alkoxy, hydrogen or nitro, in free base or acid addition salt form.

2. (Currently amended) A process for the production of a compound of formula I as defined in claim 1 and its acid addition salts, comprising the steps of

a) for the production of a compound of formula I which contains no <sup>11</sup>C or <sup>18</sup>F atom, reacting a compound of formula II



wherein X is as defined in claim 1 and Hal is Cl, Br or I, with 5-(2-fluoro-ethylamino)thiazolyl-2-boronic acid or a compound of formula III



wherein Y and R<sub>3</sub> are as defined above and R'<sub>2</sub> is a group R<sub>2</sub> as defined above which contains no <sup>11</sup>C or <sup>18</sup>F atom, or

- b) for the production of a compound of formula I wherein R<sub>1</sub> is 5-(2-<sup>18</sup>F-ethylamino)-thiazol-2-yl, reacting a compound of formula I wherein R<sub>1</sub> is 5-(2-mesyloxy-ethylamino)-thiazol-2-yl or 5-(2-tosyloxy-ethylamino)-thiazol-2-yl with <sup>18</sup>F<sup>⊖</sup>, or
- c) for the production of a compound of formula I wherein R<sub>2</sub> is NH<sup>11</sup>CH<sub>3</sub>, N(CH<sub>3</sub>)<sup>11</sup>CH<sub>3</sub> or N(<sup>11</sup>CH<sub>3</sub>)<sub>2</sub>, reacting a compound of formula I wherein R<sub>2</sub> is NH<sub>2</sub> or NHCH<sub>3</sub> with <sup>11</sup>CH<sub>3</sub>I, or
- d) for the production of a compound of formula I wherein R<sub>2</sub> is NH(CH<sub>2</sub>)<sub>n</sub><sup>18</sup>F, N(CH<sub>3</sub>)-(CH<sub>2</sub>)<sub>n</sub><sup>18</sup>F, O-(CH<sub>2</sub>)<sub>n</sub><sup>18</sup>F or CONH(CH<sub>2</sub>)<sub>n</sub><sup>18</sup>F, reacting a compound of formula I wherein R<sub>2</sub> is, respectively, NH(CH<sub>2</sub>)<sub>n</sub>OTs or NH(CH<sub>2</sub>)<sub>n</sub>OMs, N(CH<sub>3</sub>)-(CH<sub>2</sub>)<sub>n</sub>OTs or N(CH<sub>3</sub>)-(CH<sub>2</sub>)<sub>n</sub>OMs, O-(CH<sub>2</sub>)<sub>n</sub>OTs or O-(CH<sub>2</sub>)<sub>n</sub>OMs, or CONH(CH<sub>2</sub>)<sub>n</sub>OTs or ONH(CH<sub>2</sub>)<sub>n</sub>OMs, with <sup>18</sup>F<sup>⊖</sup>,

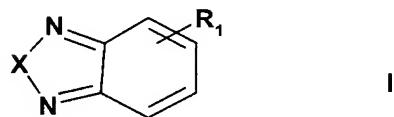
and recovering the resulting compound of formula I in free base form or in acid addition salt form ~~or an acid addition salt~~.

3. (Original) A composition for labeling histopathological structures in vitro or in vivo, comprising a compound of formula I as defined in claim 1, in free base or acid addition salt form.

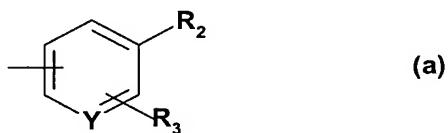
4. (Withdrawn) A method for labeling histopathological structures in vitro or in vivo, comprising contacting brain tissue with a compound of formula I as defined in claim 1, in free base or acid addition salt form.

5. (Withdrawn) A method according to claim 4, for labeling β-amyloid deposits.

6. (Withdrawn) A method according to claim 4, comprising administering the compound of formula I to a patient.
7. (Withdrawn) A method according to claim 4, comprising the further step of determining whether the compound of formula I labeled the target structure.
8. (Withdrawn) A method according to claim 7, comprising observing the target structure labeled with a non-radioactive compound of formula I, using fluorescence microscopy.
9. (Withdrawn) A method according to claim 7, comprising observing the target structure labeled with a radioactive compound of formula I, using positron emission tomography (PET).
10. (Withdrawn) A method according to claim 4 for diagnosing Alzheimer's disease.
11. (Withdrawn) A method according to claim 10, for monitoring the effectiveness of a therapeutic treatment of Alzheimer's disease.
12. (Withdrawn) A method according to claim 4, for detecting histopathological hallmarks of Alzheimer's disease.
- 13-15. (Cancelled)
16. (New) A package comprising a compound of formula I,



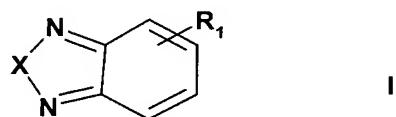
wherein X is O or S, R<sub>1</sub> is 5-(2-fluoro-ethylamino)-thiazol-2-yl, 5-(2-<sup>18</sup>F-ethylamino)-thiazol-2-yl or a group of formula (a)



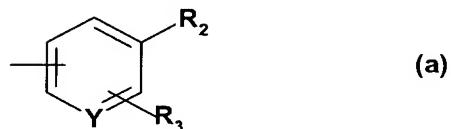
wherein Y is CH or N, R<sub>2</sub> is NH<sub>2</sub> or NHCH<sub>3</sub>, and R<sub>3</sub> is hydroxy, (C1-4)alkoxy, hydrogen or nitro, in free base or acid addition salt form,

together with instructions for the production of the compound of formula I wherein R<sub>2</sub> is NH<sup>11</sup>CH<sub>3</sub>, N(CH<sub>3</sub>)<sup>11</sup>CH<sub>3</sub> or N(<sup>11</sup>CH<sub>3</sub>)<sub>2</sub> by reaction of the starting material with freshly prepared <sup>11</sup>CH<sub>3</sub>I.

17. (New) A package comprising as starting material a compound of formula I,



wherein X is O or S, R<sub>1</sub> is 5-(2-fluoro-ethylamino)-thiazol-2-yl, 5-(2-<sup>18</sup>F-ethylamino)-thiazol-2-yl or a group of formula (a)



wherein Y is CH or N, R<sub>2</sub> is NH(CH<sub>2</sub>)<sub>n</sub>OTs, NH(CH<sub>2</sub>)<sub>n</sub>OMs, N(CH<sub>3</sub>)-(CH<sub>2</sub>)<sub>n</sub>OTs, N(CH<sub>3</sub>)-(CH<sub>2</sub>)<sub>n</sub>OMs, O-(CH<sub>2</sub>)<sub>n</sub>OTs, O-(CH<sub>2</sub>)<sub>n</sub>-OMs, CONH(CH<sub>2</sub>)<sub>n</sub>OTs or ONH(CH<sub>2</sub>)<sub>n</sub>OMs (n being in each case 2 to 4), wherein OMs corresponds to mesylate and OTs to tosylate, and R<sub>3</sub> is hydroxy, (C1-4)alkoxy, hydrogen or nitro, in free base or acid addition salt form,

together with instructions for the production of the compound of formula I wherein R<sub>2</sub> is NH(CH<sub>2</sub>)<sub>n</sub><sup>18</sup>F, N(CH<sub>3</sub>)-(CH<sub>2</sub>)<sub>n</sub><sup>18</sup>F, O-(CH<sub>2</sub>)<sub>n</sub><sup>18</sup>F or CONH(CH<sub>2</sub>)<sub>n</sub><sup>18</sup>F by a suitable reaction cascade of the starting material with <sup>18</sup>F<sup>0</sup>.